

# OPTIMA HEALTH PLAN

## PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST\*

**Directions:** The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; **1-800-750-9692**. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. If the information provided is not complete, correct, or legible, the authorization process can be delayed.

**Drug Requested:** Sucraid® (sacrosidase)

<b>DRUG INFORMATION:</b> Authorization may be delayed if incomplete.
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**Drug Form/Strength:** \_\_\_\_\_

**Dosing Schedule:** \_\_\_\_\_ **Length of Therapy:** \_\_\_\_\_

**Diagnosis:** \_\_\_\_\_ **ICD Code, if applicable:** \_\_\_\_\_

**Quantity Limit:** 236mL/30 days

<b>CLINICAL CRITERIA:</b> Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.
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<b><u>Initial Approval:</u> 60 days</b>
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- ☐ Patient is 5 months of age or older and has a diagnosis of congenital sucrase-isomaltase deficiency (CSID) confirmed by a gastroenterologist, endocrinologist, or genetics specialist

**AND**

- ☐ Patient has documented chronic symptoms of CSID including watery diarrhea, abdominal pain, gas/bloating after sucrose/starch ingestion (must submit chart notes documenting symptoms following sucrose/starch ingestion)
- Number of severe GI events within the last 2 months: \_\_\_\_\_ (must be documented in submitted chart notes)

**AND**

- ☐ A low sucrose and low starch diet has been attempted with improvement in patient symptoms, and patient will continue to follow a low sucrose, low starch diet while on therapy

**AND**

- ☐ Patient does not have lactose intolerance or a secondary sucrase deficiency associated with any of the following: celiac disease, Crohn's disease, autoimmune gastroenteropathy, eosinophilic gastroenteropathy, short bowel syndrome, Giardiasis, small intestinal bacterial overgrowth (SIBO), acute gastroenteritis, or enteropathy associated with acquired immune deficiency syndrome

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AND (**ALL** 4 below **MUST** be met):

<input type="checkbox"/> Stool pH < 6.0	<input type="checkbox"/> Increase in breath hydrogen of > 10 ppm when challenged with sucrose after fasting
<input type="checkbox"/> Genetic test results confirm diagnosis of CSID	<input type="checkbox"/> Negative lactose breath test

**OR (BOTH below MUST be met)**

- ☐ Small bowel biopsy documents intestinal sucrase activity of <25 U/g protein (must be greater than 2 standard deviations below the mean) with normal or decreased maltase and isomaltase levels, normal levels of other disaccharides, and normal villous architecture of the small intestine on biopsy
- ☐ Genetic testing results document sucrase-isomaltase deficiency (CSID)

**Reauthorization Approval: 12 months.** Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

- ☐ Patient has had a 50% reduction in all symptoms of CSID, including watery diarrhea, abdominal pain, gas/bloating; etc. (**improvement from baseline must be noted in submitted chart notes**)
  - Number of severe GI events within the last 2 months: \_\_\_\_\_ (**must be documented in submitted chart notes**)

AND

- ☐ Patient will continue to follow a low sucrose, low starch diet while on therapy

**Medication being provided by Specialty Pharmacy - PropriumRx**

*Not all drugs may be covered under every Plan*

*If a drug is non-formulary on a Plan, documentation of medical necessity will be required.*

***\*\*Use of samples to initiate therapy does not meet step edit/ preauthorization criteria.\*\****

***\*Previous therapies will be verified through pharmacy paid claims or submitted chart notes.\****

Patient Name: \_\_\_\_\_

Member Optima #: \_\_\_\_\_ Date of Birth: \_\_\_\_\_

Prescriber Name: \_\_\_\_\_

Prescriber Signature: \_\_\_\_\_ Date: \_\_\_\_\_

Office Contact Name: \_\_\_\_\_

Phone Number: \_\_\_\_\_ Fax Number: \_\_\_\_\_

DEA OR NPI #: \_\_\_\_\_

\*Approved by Pharmacy and Therapeutics Committee: 4/15/2015; 8/20/2020

REVISED/UPDATED: 1/26/2015; 5/22/2015; 12/29/2015; 12/20/2016; 8/18/2017; 12/30/2018; (Reformatted) 6/19/2019; 11/12/2020